

Serial No. 09/411,568
STN SEARCH

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=> file

ENTER A FILE NAME OR (HOME):caplus

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FULL ESTIMATED COST 0.45 0.45
FILE 'CAPLUS' ENTERED AT 10:12:25 ON 12 DEC 2000
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FILE COVERS 1967 - 12 Dec 2000 VOL 133 ISS 25
FILE LAST UPDATED: 11 Dec 2000 (20001211/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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Serial No. 09/411,568
STN SEARCH

Now you can extend your author, patent assignee, patent information, and title searches back to 1907. The records from 1907-1966 now have this searchable data in CAOLD. You now have electronic access to all of CA: 1907 to 1966 in CAOLD and 1967 to the present in CAPLUS on STN.

=> s (itr or (internal (w) tandem (w) repeat#))/bi,ab

385 ITR/BI
319 ITR/AB
235311 INTERNAL/BI
209767 INTERNAL/AB
27102 TANDEM/BI
23200 TANDEM/AB
48590 REPEAT#/BI
44412 REPEAT#/AB
14 INTERNAL (W) TANDEM (W) REPEAT#
L1 398 (ITR OR (INTERNAL (W) TANDEM (W) REPEAT#))/BI,AB

=> s ((anti (w) sense) or antisense)/bi,ab

223038 ANTI/BI
187535 ANTI/AB
20686 SENSE/BI
19836 SENSE/AB
849 ANTI (W) SENSE
17671 ANTISENSE/BI
14720 ANTISENSE/AB
L2 18196 ((ANTI (W) SENSE) OR ANTISENSE)/BI,AB

=> s l1 and l2

L3 5 L1 AND L2

=> s ribozyme#/bi,ab

4477 RIBOZYME#/BI
3235 RIBOZYME#/AB
L4 4477 RIBOZYME#/BI,AB

=> s l3 and l4

L5 2 L3 AND L4

=> d l3 1-5 bib ab

L3 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2000 ACS
AN 1999:795943 CAPLUS
DN 132:45813
TI Generation of recombinant adeno-associated virus vectors without formation of wild-type virus
IN Srivastava, Arun; Wang, Xu-Shan; Ponnazhagan, Selvarangan PA Advanced Research and Technology Institute, USA
SO PCT Int. Appl., 100 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9964569 A1 19991216 WO 1999-US13070 19990609 W:
AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
AU 9945587 A1 19991230 AU 1999-45587 19990609 PRAI US 1998-88714 19980610
WO 1999-US13070 19990609

AB A plasmid co-transfection system for the generation of recombinant adeno-assocd. virus 2 for use as a gene delivery virus that minimizes the generation of wild-type virus by preventing homologous recombination is described. Recombination is dependent upon 10 nucleotides of the viral D-sequence and helper vectors lacking sequence homol. in the D-sequence and helper plasmids lacking adenovirus inverted terminal repeats. Methods and compns. for the use of recombinant AAV plasmids and helper vectors lacking homol. in the D-sequence, and helper plasmids lacking the adenovirus ITRs for use in gene therapy are described. Mapping of recombination events leading to the generation of wild-type virus found most of them clustering in the 10 distal nucleotides of the D-sequence and also involved the inverted terminal repeats of the adenovirus 5 helper. Deletion of selected sequences gradually lowered the titer of wild-type virus to <0.1% of total virus.

RE.CNT 6

RE

- (1) Qing; Journal of Virology 1998, V72(2), P1593 CAPLUS
- (2) Wang; Journal of Molecular Biology 1995, V250, P573 CAPLUS (3) Wang; Journal of Virology 1996, V70(3), P1668 CAPLUS
- (4) Wang; Journal of Virology 1997, V71(2), P1140 CAPLUS
- (5) Wang; Journal of Virology 1997, V71(4), P3077 CAPLUS

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2000 ACS
AN 1999:244775 CAPLUS
DN 130:292438
TI Chimeric AAV/B19 parvovirus-based recombinant vector system specifically targeting the erythroid lineage
IN Srivastava, Arun; Ponnazhagan, Selvarangan PA Advanced Research and Technology Institute, USA
SO PCT Int. Appl., 76 pp.
CODEN: PIXXD2
DT Patent

Serial No. 09/411,568
STN SEARCH

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9918227 A1 19990415 WO 1998-US21202 19981008 W:

AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN,
CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU,
ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA,
UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU,
TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE,
CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML,
MR, NE, SN, TD, TG

AU 9912696 A1 19990427 AU 1999-12696 19981008 EP
1027451 A1 20000816 EP 1998-956097 19981008 R: AT, BE,
CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
PT, IE, FI

PRAI US 1997-61364 19971008

WO 1998-US21202 19981008

AB The present invention relates to the engineering,
propagation and use of chimeric parvovirus vectors
using sequences from adeno-assocd. virus (AAV) and B19
virus, which may be used to deliver genes to various
target cells, including those of erythroid lineage.
The system exploits the unique features of AAV and B19
such that it does not suffer from toxicity,
oncogenicity, or immunogenicity concerns. Heterologous
DNA sequences are cloned withing the inverted terminal
repeats (ITR) of AAV, without the presence of any
AAV structural genes, and subsequently packaged inside
the capsid structure of B19. Such a chimeric vector is
achieved by creating a helper plasmid consisting of
the rep gene of AAV, and the cap gene of B19. High
titers of the vector may be generated, facilitating in
vivo therapy. It is designed to specifically target
primitive progenitor and differentiated cells of
erythroid lineage, and can achieve stable integration
and expression of transduced genes. RE.CNT 11
RE

(1) Childrens Hospital Inc; WO 9534670 A 1995 CAPLUS

(2) Latta, M; WO 9523867 A 1995 CAPLUS

(4) Ponnazhagan, S; Blood, Meeting Info: 39th Annual
Meeting of the American Society of Hematology 1997
CAPLUS

(5) Ponnazhagan, S; J Virology 1998, V72(6), P5224
CAPLUS

(7) RES Corp Technologies Inc; WO 9309239 A 1993
CAPLUS

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2000 ACS

AN 1998:303188 CAPLUS

DN 129:77198

TI Site-specific integration in mammalian cells
mediated by a new hybrid baculovirus-adeno-associated
virus vector

AU Palombo, Fabio; Monciotti, Andrea; Recchia,
Alessandra; Cortese, Riccardo; Ciliberto, Gennaro; La
Monica, Nicola

CS IRBM P. Angeletti, Pomezia, 00040, Italy

SO J. Virol. (1998), 72(6), 5025-5034

CODEN: JOVIAM; ISSN: 0022-538X

PB American Society for Microbiology

DT Journal

LA English

AB Baculovirus can transiently transduce primary human
and rat hepatocytes, as well as a subset of stable
cell lines. To prolong transgene expression, we have
developed new hybrid vectors which assoc. key elements
from adeno-assocd. virus (AAV) with the elevated
transducing capacity of baculovirus. The hybrid
vectors contain a transgene cassette composed of the
.beta.-galactosidase (.beta.-Gal) reporter gene and
the hygromycin resistance (Hygr) gene flanked by the
AAV inverted terminal repeats (ITRs), which are
necessary for AAV replication and integration in the
host genome. Constructs were derived both with and
without the AAV rep gene under the p5 and p19
promoters cloned in different positions with respect
to the baculovirus polyhedrin promoter. A high-titer
prepn. of baculovirus-AAV (Bac-AAV) chimeric virus
contg. the ITR -Hygr-.beta.-Gal sequence was obtained
with insect cells only when the rep gene was placed in
an antisense orientation to the polyhedrin promoter.
Infection of 293 cells with Bac-AAV virus expressing
the rep gene results in a 10-to 50-fold increase in
the no. of Hygr stable cell clones. Addnl., rep
expression detd. the localization of the transgene
cassette in the aavsl site in approx. 41% of cases as
detected by both Southern blotting and fluorescent in
situ hybridization anal. Moreover, site-specific
integration of the ITR -flanked DNA was also detected
by PCR amplification of the ITR -aavsl junction in
transduced human fibro-blasts. These data indicate
that Bac-AAV hybrid vectors can allow permanent,
nontoxic gene delivery of DNA constructs for ex vivo
treatment of primary human cells.

L3 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2000 ACS

AN 1998:169418 CAPLUS

DN 128:227084

TI Methods and compositions for liver-specific
delivery of therapeutic molecules using recombinant
adeno-associated virus vectors IN Srivastava, Aron;
Ponnazhagan, Selvarangan; Chloemer, Robert H.; Wang,
Xu-Shan; Yoder, Mervin C.; Zhou, Shang-Zhen; Escobedo,
Jaime; Dwarkl, Varavani

PA Chiron Corporation, USA; Indiana University

SO PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

Serial No. 09/411,568
STN SEARCH

PI WO 9809524 A1 19980312 WO 1997-US15453 19970902 W:
CA, JP

RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT,
LU, MC, NL, PT, SE EP 933997 A1 19990811 EP
1997-940762 19970902 R: AT, BE, CH, DE, DK, ES, FR,
GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI
PRAI US 1996-25616 19960906

US 1996-25649 19960911

WO 1997-US15453 19970902

AB Provided are methods for selectively expressing
therapeutic mols., such as secretory proteins,
antisense mols. and ribozymes, in the liver. The
methods find use in treating hepatic diseases or
conditions. The methods also find use in treating any
disease or condition in which systemic administration
of the therapeutic substance, for example, a secretory
protein, is desired. The methods involve administering
to a mammalian patient having a need for liver
expression of a therapeutic mol. an AAV vector contg.
a therapeutically effective amt. of the therapeutic
mol. Also provided are novel vectors employable in
these methods. Expts. revealed that, following i.v.
injection of AAV vectors into mice, the AAV genomes
were found predominantly in the liver. The
heterologous genes carried by these vectors (chimeric
cytomegalovirus promoter-lacZ or .beta.-globin
promoter-globin genes) were expressed in the liver.
Cotransfection of adenovirus 2-infected 293 cells with
the AAV vectors and helper plasmid contg. cap and rep
genes resulted in prodn. of 0.1-10% wild-type AAV.
Replacement of the last 10 nucleotides of the ITR D
sequence with unrelated nucleotides reduced this
illegitimate recombination was reduced. Four
recombinant AAV vectors (pD-5, pD-10, pD-15 and pD-20)
with such modified ITR regions were prepd.
L3 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2000 ACS

AN 1995:951301 CAPLUS

DN 123:332111

TI Integrative adenovirus expression vectors for use
in gene therapy IN Deneffe, Patrice; Latta, Martine;
Perricaudet, Michel; Vigne, Emmanuelle PA
Rhone-Poulenc Rorer S.A., Fr.

SO PCT Int. Appl., 49 pp.

CODEN: PIXXD2

DT Patent

LA French

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9523867 A1 19950908 WO 1995-FR233 19950228 W:

AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU,
JP, KE, KG, KP, KR, KZ, LK, LR, LT, LV, MD, MG, MN,
MW, MX, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA,
US, UZ, VN

RW: KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,
CI, CM, GA, GN, ML, MR, NE, SN, TD, TG

FR 2716893 A1 19950908 FR 1994-2445 19940303 FR
2716893 B1 19960412

CA 2184113 AA 19950908 CA 1995-2184113 19950228 AU
9518526 A1 19950918 AU 1995-18526 19950228 EP 748385
A1 19961218 EP 1995-910605 19950228 R: AT, BE, CH, DE,
DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE JP
09509578 T2 19970930 JP 1995-522730 19950228 ZA
9501803 A 19960109 ZA 1995-1803 19950303 US 6033885 A
20000307 US 1996-702573 19960912 PRAI FR 1994-2445
19940303

WO 1995-FR233 19950228

AB Recombination-defective adenoviruses carrying a
cassette that can be integrated into the genome of
host cells are constructed for use in gene therapy.
The cassette particularly contains at least one
inverted terminal repeat (ITR) of an adeno-assocd.
virus (AAV) and a therapeutic gene. The use of the AAV
ITR directs integration to the same locus in all cases
and minimizes possible complications from random
integration. The construction of virus carrying the
lacZ reporter gene or a human lipoprotein AI gene
under control of viral (vesicular stomatitis or Rous
sarcoma virus) promoters is described.

=> file ca

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FILE COVERS 1967 - 7 Dec 2000 VOL 133 ISS 25

FILE LAST UPDATED: 7 Dec 2000 (20001207/ED)

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of CA: 1907 to 1966 in CAOLD and 1967 to the present
in CA on STN.

=> s 15

374 ITR/BI

309 ITR/AB

Serial No. 09/411,568
STN SEARCH

228219 INTERNAL/BI
203130 INTERNAL/AB
25935 TANDEM/BI
22201 TANDEM/AB
46369 REPEAT#/BI
42301 REPEAT#/AB
13 INTERNAL (W) TANDEM (W) REPEAT#
215262 ANTI/BI
180573 ANTI/AB
19151 SENSE/BI
18427 SENSE/AB
813 ANTI (W) SENSE
16873 ANTISENSE/BI
14081 ANTISENSE/AB
4250 RIBOZYME#/BI
3054 RIBOZYME#/AB
L6 2 L3 AND L4

=> log y

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE
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STN INTERNATIONAL LOGOFF AT 10:14:25 ON 12 DEC 2000